California Drug Recall Information

Recall Name
Hospira Recalls Lidocaine HCl Injection Due to Particulate Matter

Recall Date	Product Description	Recalling Firm	Recall Reason
12/23/13	Lidocaine HCl Injection, USP, 2% 5 ml single-dose vial **NDC # 0409-2066-05**	**Hospira, Inc.** Lake Forest, IL	*Due to a reddish orange particulate on the inner surface and floating in the solution.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
N/A	**Lot 32-135-DD** **Expires 1AUG2015** Product Labels	**CA**, nationwide	Distributed between: September 2013 and October 2013.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm379739.htm